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EXAMINER

SAVAGE, MATTHEW O

ART UNIT

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Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

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The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negated by the manner in which the invention was made.

Claims 1, 3, 4, and 6-14 are rejected under 35 U.S.C. 103(a) as being unpatentable over Brym in view of Zhang et al.

With respect to claim 1, Brym discloses a method for removing mercury from a fluid stream (see line 66 of col. 6, the fluid being water as mentioned in the abstract) including the steps of: providing a composite material comprising a substrate (e.g., plates of glass or ceramic, see lines 48-49 of col. 8) and catalyst particles (e.g., TiO_2 , see lines 50-51 of col. 8); and contacting a fluid stream with the composite, wherein the composite adsorbs and oxidizes the mercury (e.g., to MgO , see line 66 of col. 6, the TiO_2 being capable of absorbing and oxidizing the mercury). Brym fails to specify the catalyst particles as being dispersed in the substrate. Zhang et al discloses a composite material that includes catalyst particles disposed within the substrate (e.g., silica gel treated by the process described on lines 32-51 of col. 10) and suggests that such a support provides a large surface area for adsorption and oxidation of oxidizing contaminants. It would have been obvious to have modified the method of Brym so as to have included the substrate in the form of a sorbent as suggested by Zhang et al in

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order to increase the amount of surface area for the adsorption and oxidation of contaminants including organic contaminants and mercury.

With respect to claim 3, Brym and Zhang et al disclose TiO_2 which can function as a sorbent. In addition, Zhang et al disclose silica gel which can function as a sorbent.

Concerning claim 4, Zhang et al disclose a gel (e.g., silica gel, see line 12 of col. 9).

Concerning claim 6, Brym and Zhang et al disclose the step of irradiating the composite material with radiation (e.g., with ultraviolet light, see lines 33-38 of col. 9 of Brym, and from line 45 of col. 5 to line 9 of col. 6 of Zhang et al).

As to claim 7, Brym discloses the radiation as having a wavelength of from about 160 to about 680 nm (e.g., 320-400 nm, see lines 36-38 of col. 9).

Concerning claim 8, Zhang et al disclose a silica gel substrate that is transparent to radiation (see lines 12-14 of col. 9).

Concerning claim 9, Zhang et al specify a substrate formed of porous silica (e.g., DavissilTM).

Regarding claim 10, Brym and Zhang et al disclose a catalyst comprising TiO_2 (see line 23 of col. 8 of Brym and line 50 of col. 10 of Zhang et al).

As to claim 11, Zhang et al discloses a support formed of silica gel, specifically DavissilTM which has a surface area of 1-1500 m^2/g .

Regarding claim 12, Brym fails to specify the catalyst as being present in an amount from .1-100%. Zhang et al teach using catalyst present in a composite material in an amount of .1-50% and teaches that such an arrangement increases the surface

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area of a catalyst bed when using the catalyst in particulate form. It would have been obvious to have modified the method of Brym so as to have included the arrangement of Zhang et al in order to increase the surface area of the catalyst bed.

Concerning claim 13, Brym fails to specify regenerating the composite. Zhang et al disclose regenerating an analogous composite (see lines 5-7 of col. 12) and suggests that such a step removes accumulated contaminants from the composite thereby maintaining the contaminate removal efficiency at a high level. It would have been obvious to have modified the method of Brym so as to have included the regeneration step as suggested by Zhang et al in order to maintain the contaminate removal efficiency of the composite at a high level.

Regarding claim 14, Zhang et al disclose thermal regeneration (e.g., with hot water or steam, see line 6 of col. 12).

Claim 5 is rejected under 35 U.S.C. 103(a) as being unpatentable over Brym in view of Zhang et al as applied to claim 4 above, and further in view of Burns et al.

Zhang et al disclose silica gel but fails to specify xerogel. Burns et al disclose xerogel and suggest that such an adsorbent has a high surface area. It would have been obvious to have modified the silica gel substrate suggested by Brym and Zang et al so as to have included a silica xerogel as suggested by Burns et al in order to further increase the surface area of the substrate thereby increasing the reactive surface area of the composite.

Applicant's arguments filed 12-12-08 have been fully considered but they are not persuasive.

Applicant argues that Zhang et al fails to disclose catalyst particles dispersed in the substrate, however, it is held that Zhang et al disclose a process that produces TiO₂ photocatalyst particles dispersed in a silica gel substrate (see from lines 32-51 of col. 10).

THIS ACTION IS MADE FINAL. Applicant is reminded of the extension of time policy as set forth in 37 CFR 1.136(a).

A shortened statutory period for reply to this final action is set to expire THREE MONTHS from the mailing date of this action. In the event a first reply is filed within TWO MONTHS of the mailing date of this final action and the advisory action is not mailed until after the end of the THREE-MONTH shortened statutory period, then the shortened statutory period will expire on the date the advisory action is mailed, and any extension fee pursuant to 37 CFR 1.136(a) will be calculated from the mailing date of the advisory action. In no event, however, will the statutory period for reply expire later than SIX MONTHS from the mailing date of this final action.

/Matthew O Savage/
Primary Examiner, Art Unit 1797
571-272-1146